Study: CDISCPILOT01 Subject: 01-701-1023 (1023) Status: ADVERSE EVENT Page 1 of 2 First Dose Date: 2012-08-05 Current Visit: RETRIEVAL

Demographics									
Age (y)	Sex Race		Ethnicity	Country	Planned Arm	Actual Arm			
64	M	WHITE	HISPANIC OR LATINO	USA	Placebo	Placebo			

Subject Visits							
Visit	Visit Day	Visit Date					
SCREENING 1	-7	2012-07-22					
SCREENING 2	-1	2012-08-03					
BASELINE	1	2012-08-05					
AMBUL ECG PLACEMENT	13	2012-08-26					
WEEK 2	14	2012-08-27					
WEEK 4	28	2012-09-02					
UNSCHEDULED 5.1		2013-02-18					
AE FOLLOW-UP	•	2013-02-18					
RETRIEVAL	168	2013-02-18					

Adverse Events									
Sponsor Defined Identifier and Verbatim Term	Start Date (Day)	End Date (Day)	Ser-ious?	Duration (days)	Sever- ity	Relation- ship	Outcome	Action Taken	Treatment Emergent?
E08: ERYTHEMA	2012-08-07	2012-08-30	No	24	Mild	Possible	Not Recovered/Not Resolved		Yes
E08: ERYTHEMA	2012-08-07	2012-08-30	No	24	Mild	Possible	Recovered/Resolved		Yes
E09: ERYTHEMA	2012-08-07		No		Moderate	Probable	Not Recovered/Not Resolved		Yes
E10: ATRIOVENTRICULAR BLOCK SECOND DEGREE	2012-08-26		No		Mild	Possible	Not Recovered/Not Resolved		Yes

Study: CDISCPILOT01 Page 2 of 2
Subject: 01-701-1023 (1023) First Dose Date: 2012-08-05
Status: ADVERSE EVENT Current Visit: RETRIEVAL

Vital Signs								
Vital Sign Parameter	SCR1	SCR2	BSLN	ECPL	WK02	WK04	RETR	
Diastolic BP (mmHg)	90	85	88	89	90	92	84	
Pulse Rate (BEATS/MIN)	74	92	76	77	94	70	76	
Systolic BP (mmHg)	140	138	132	138	138	136	130	
Temperature (C)	36.6	36.6	36.3	36.4	36.9	36.2	36.6	
Weight (kg)	78.5		80.3		80.7	80.3		

